

IN THE CLAIMS:

Please amend claims 1, 53, 59, 63 and 66 as follows. Please cancel claims 2 and 56 without prejudice or disclaimer. This listing of claims will replace all prior versions, and listings of claims, in the application.

LISTING OF CLAIMS:

1. (Currently amended) A method of producing and identifying a mammalian protease mutein that inactivates an activity of a target protein involved with a disease or pathology in a mammal, wherein:

~~the target protein is selected from among a cell surface molecule that transmits an extracellular signal for cell proliferation, a cytokine, a cytokine receptor and a signaling protein that regulates apoptosis;~~

~~inactivation of the target protein can ameliorate a disease or pathology;~~

~~cleavage of a substrate sequence in the target protein inactivates an activity of the target protein; and~~

~~the method comprises~~ comprising the steps of:

(a) producing a library of comprising protease muteins comprising mutation(s) of a serine protease scaffold and/or biologically active portions thereof at an amino acid position selected from among amino acid residues between 58 and 64, 97, 98, 99, 100, 171, 174, 180, 189, 190, 191, 192, 215, 217, 218 and 226 by chymotrypsin numbering, wherein:

each different mutein protease in the library is a member of the library;

each member of the library has N mutations relative to a wild-type mammalian protease scaffold or a biologically active portion thereof; and

N is a positive integer;

(b) contacting members of the library with ~~the~~ a target protein or with a polypeptide comprising a substrate sequence that is present in the target protein, wherein:

the target protein is selected from among a cell surface molecule that transmits an extracellular signal for cell proliferation, a cytokine, a cytokine receptor and a signaling protein that regulates apoptosis; and

cleavage of a substrate sequence in the target protein inactivates an activity of the target protein; and

(c) measuring a cleavage activity and/or substrate specificity of at least two members of the library for the target protein or substrate sequence;

(d) based on the measured activity and/or specificity, identifying members of the library that have an increased cleavage activity and/or altered substrate specificity for cleaving the substrate sequence or the target protein, relative to the wild-type mammalian protease scaffold; and

(e) testing the identified protease(s) or biologically active portion thereof for cleavage and inactivation of an activity of the target protein that contains the substrate sequence, thereby identifying a protease mutein or a biologically active portion thereof that inactivates an activity of a target protein that is involved with the disease or pathology.

2. (Cancelled)
3. (Previously presented) The method of claim 1, wherein N is an integer between 1 and 20.
4. (Previously presented) The method of claim 3, wherein N is an integer from 1-5.
5. (Previously presented) The method of claim 3, wherein N is an integer from 5-10.
6. (Previously presented) The method of claim 3, wherein N is an integer from 10-20.
7. (Previously presented) The method of claim 1, wherein the wild-type mammalian protease is selected from among trypsin, chymotrypsin, subtilisin, MTSP-1, granzyme A, granzyme B, and granzyme M, elastase, chymase, papain, neutrophil elastase, complement factor serine proteases, ADAMTS13, neural endopeptidase/neprilysin, furin, cruzain, and urokinase plasminogen activator (uPA).
8. (Cancelled)
9. (Previously presented) The method of claim 1, wherein the pathology is selected from among rheumatoid arthritis, sepsis, cancer, acquired immunodeficiency syndrome, respiratory tract infections, influenza, cardiovascular disease and asthma.
10. (Canceled)
11. (Original) The method of claim 1, wherein the target protein is involved in apoptosis.
12. (Previously presented) The method of claim 11, wherein the target protein is caspase-3, VEGF or VEGF-R.
13. (Previously presented) The method of claim 1, wherein the specificity of the identified protease mutein for cleaving the substrate sequence is increased by at least 10-fold

compared to the specificity of the wild-type mammalian protease scaffold for cleaving the substrate sequence.

14. (Previously presented) The method of claim 1, wherein the specificity of the identified protease mutein for cleaving the substrate sequence is increased by at least 100-fold compared to the specificity of the wild-type mammalian protease scaffold for cleaving the substrate sequence.

15. (Previously presented) The method of claim 1, wherein the specificity of the identified protease mutein for cleaving the substrate sequence is increased by at least 1000-fold compared to the specificity of the wild-type mammalian protease scaffold for cleaving the substrate sequence.

16. (Previously presented) The method of claim 1, further comprising the steps of:

(e) identifying the mutation(s) contained in a first mutein protease and a second mutein protease identified in step c) as having increased cleavage activity and/or altered specificity;

(f) producing a third mutein protease containing the mutations of the first mutein protease and the mutations of the second mutein protease; and

(g) measuring the cleavage activity and/or substrate specificity of the third mutein protease to determine its cleavage activity and/or specificity for the substrate sequence compared to the first mutein protease or second mutein protease.

17-44. (Canceled)

45. (Previously presented) The method of claim 1, further comprising:
repeating steps a)-d) iteratively to produce a further library of protease muteins from the identified muteins each with increased cleavage activity and/or altered substrate specificity; and

identifying among the further library, a mutein protease having altered specificity and/or increased cleavage activity relative to the wild-type mammalian protease.

46. (Cancelled)

47. (Cancelled)

48. (Previously presented) The method of claim 1, wherein the substrate sequence is a sequence in a target protein that is a human protein.

49.-50. (Cancelled)

51. (Previously presented) The method of claim 50, wherein the member of the library identified in step (d) has the highest measured cleavage activity for the substrate sequence among the identified members of the library.

52. (Previously presented) The method of claim 1, further comprising the steps of providing at least one mutein protease identified in step (d), and testing the mutein protease in an *in vivo* assay for inactivation of an activity of the target protein.

53. (Currently amended) A method of producing and identifying a mammalian protease mutein that inactivates an activity of a target protein involved with a disease or pathology in a mammal, ~~wherein:~~

~~inactivation of the target protein can ameliorate a disease or pathology;~~

~~cleavage of a substrate sequence in the target protein inactivates an activity of the target protein;~~

~~the mammalian protease is selected from among a granzyme A, granzyme B, granzyme M, cathepsin, trypsin, chymotrypsin, subtilisin, MTSP-1, elastase, chymase, tryptase, collagenase, papain, neutrophil elastase, complement factor serine proteases, ADAMTS13, neural endopeptidase/neprilysin, furin, cruzain, and urokinase plasminogen activator (uPA); and~~

~~the method comprises~~ comprising the steps of:

(a) producing a library ~~of comprising~~ comprising protease muteins comprising mutation(s) of a mammalian protease scaffold and/or biologically active portion thereof at an amino acid position selected from among amino acid residues between 58 and 64, 97, 98, 99, 100, 171, 174, 180, 189, 190, 191, 192, 215, 217, 218 and 226 by chymotrypsin numbering, wherein:

the mammalian protease scaffold is selected from among a granzyme A, granzyme B, granzyme M, cathepsin, trypsin, chymotrypsin, subtilisin, MTSP-1, elastase, chymase, tryptase, collagenase, papain, neutrophil elastase, complement factor serine proteases, ADAMTS13, neural endopeptidase/neprilysin, furin, cruzain, and urokinase plasminogen activator (uPA);

each different protease mutein in the library being a member of the library;

each member of the library has N mutations relative to a wild-type mammalian protease scaffold or a biologically active portion thereof; and

N is a positive integer;

(b) contacting members of the library with ~~the a~~ a target protein or with a polypeptide comprising a substrate sequence that is present in the target protein, wherein:

the target protein is involved with a disease or pathology in a mammal; and
cleavage of a substrate sequence in the target protein inactivates an activity of
the target protein;

(c) measuring a cleavage activity and/or substrate specificity of at least two members of the library for the target protein or substrate sequence;

(d) based on the measured activity and/or specificity, identifying those members of the library that have an increased cleavage activity and/or altered substrate specificity for cleaving the substrate sequence or the target protein relative to the wild-type mammalian protease scaffold;

(e) testing the identified mutein protease(s) or biologically active portion thereof for cleavage and inactivation of an activity of the target protein that contains the substrate sequence, thereby identifying a protease mutein or a biologically active portion thereof that inactivates an activity of a target protein that is involved with or causes the disease or pathology;

(f) identifying the mutation(s) contained in a first mutein protease and a second mutein protease identified in step (d) step (e) as having increased cleavage activity and/or altered specificity for cleaving the target protein or substrate sequence and that can inactivate an activity of the target protein;

(g) generating a third mutein protease containing the mutations of the first mutein protease and the mutations of the second mutein; and

(h) measuring the cleavage activity and/or substrate specificity of the third mutein protease to determine whether the third mutein protease has increased cleavage activity and/or altered specificity toward the target protein or substrate sequence compared to the first mutein protease or second mutein protease; and

(i) testing the third mutein for cleavage and inactivation of an activity of the target protein that contains the substrate sequence, thereby identifying a protease mutein or a biologically active portion thereof that inactivates an activity of a target protein that is involved with or causes the disease or pathology.

54. (Cancelled)

55. (Cancelled)

56. (Cancelled)

57. (Previously presented) The method of claim 53, further comprising:

repeating steps a)-d) iteratively to produce a further library of protease muteins from the identified muteins each with increased cleavage activity and/or altered substrate specificity; and

identifying among the further library, a mutein protease having altered specificity and/or increased cleavage activity for the substrate sequence relative to the wild-type mammalian protease.

58. (Previously presented) The method of claim 53, wherein the substrate sequence is a sequence in a target protein that is a human protein.

59. (Currently amended) A method of producing and identifying a human protease mutein that inactivates an activity of a target protein involved with a disease or pathology in a human, ~~wherein:~~

~~— cleavage of a substrate sequence in the target protein inactivates an activity of the target protein;~~

~~— the target protein is selected from among a cell surface molecule that transmits an extracellular signal for cell proliferation, a cytokine, a cytokine receptor and a signaling protein that regulates apoptosis;~~

~~— inactivation of the target protein can ameliorate a disease or pathology; and~~

~~— the method comprises~~ comprising the steps of:

(a) producing a library ~~of comprising~~ protease muteins comprising mutation(s) of a human protease scaffold and/or biologically active portions thereof at an amino acid position selected from among amino acid residues between 58 and 64, 97, 98, 99, 100, 171, 174, 180, 189, 190, 191, 192, 215, 217, 218 and 226 by chymotrypsin numbering, wherein:

each different protease mutein in the library is a member of the library;

each member having N mutations relative to a wild-type human protease scaffold or a biologically active portion thereof, wherein N is a positive integer; and

the human protease scaffold is selected from among a granzyme A, granzyme B, granzyme M, cathepsin, trypsin, chymotrypsin, subtilisin, MTSP-1, elastase, chymase, tryptase, collagenase, papain, neutrophil elastase, complement factor serine proteases, ADAMTS13, neural endopeptidase/neprilysin, furin, cruzain, and urokinase plasminogen activator (uPA);

(b) contacting members of the library with the target protein or with a polypeptide comprising a substrate sequence that is present in the target protein, wherein:

the target protein is selected from among a cell surface molecule that transmits an extracellular signal for cell proliferation, a cytokine, a cytokine receptor and a signaling protein that regulates apoptosis; and

cleavage of a substrate sequence in the target protein inactivates an activity of the target protein;

(c) measuring a cleavage activity and/or substrate specificity of at least two members of the library for the substrate sequence or the target protein; and

(d) based on the measured activity and/or specificity, identifying those members of the library that have an increased cleavage activity and/or altered substrate specificity for cleaving the substrate sequence or the target protein relative to the wild-type human protease scaffold; and

(e) testing the identified protease(s) for cleavage and inactivation of an activity of the target protein that contains the substrate sequence, thereby identifying a protease mutein or a biologically active portion thereof that inactivates an activity of a target protein that is involved with or causes the disease or pathology.

60. (Cancelled)

61. (Previously presented) The method of claim 59, wherein the wild-type protease scaffold is Granzyme B or MTSP-1.

62. (Previously presented) The method of claim 59, wherein the target protein is selected from among caspase 3, tumor necrosis factor, tumor necrosis factor receptor, interleukin-1, interleukin-1 receptor, interleukin-2, interleukin-2 receptor, interleukin-4, interleukin-4 receptor, interleukin-5, interleukin-5 receptor, interleukin-12, interleukin-12 receptor, interleukin-13, interleukin-13 receptor, p-selectin, p-selectin glycoprotein ligand, Substance P, Bradykinin, PSGL, factor IX, immunoglobulin E, immunoglobulin E receptor, CCR5, CXCR4, glycoprotein 120, glycoprotein 41, hemagglutinin, respiratory syncytium virus fusion protein, B7, CD28, CD2, CD3, CD4, CD40, vascular endothelial growth factor, VEGF receptor, fibroblast growth factor, endothelial growth factor, EGF receptor, TGF receptor, transforming growth factor, Her2, CCR1, CXCR3, CCR2, Src, Akt, Bcl-2, BCR-Abl, glucagon synthase kinase-3, cyclin dependent kinase-2 (cdk-2) and cyclin dependent kinase-4 (cdk-4).

63. (Currently amended) A method of producing and identifying a human protease mutein that inactivates an activity of a target protein involved with a disease or pathology in a mammal, ~~wherein:~~

~~the cleavage of a substrate sequence in the target protein inactivates an activity of the target protein;~~

~~the target protein is selected from among a caspase 3, tumor necrosis factor, tumor necrosis factor receptor, interleukin 1, interleukin 1 receptor, interleukin 2, interleukin 2 receptor, interleukin 4, interleukin 4 receptor, interleukin 5, interleukin 5 receptor, interleukin 12, interleukin 12 receptor, interleukin 13, interleukin 13 receptor, p-selectin, p-selectin glycoprotein ligand, Substance P, Bradykinin, PSGL, factor IX, immunoglobulin E, immunoglobulin E receptor, CCR5, CXCR4, glycoprotein 120, glycoprotein 41, hemagglutinin, respiratory syncytium virus fusion protein, B7, CD28, CD2, CD3, CD4, CD40, vascular endothelial growth factor, VEGF receptor, fibroblast growth factor, endothelial growth factor, EGF receptor, TGF receptor, transforming growth factor, Her2, CCR1, CXCR3, CCR2, Src, Akt, Bel 2, BCR-Abl, glucagon synthase kinase 3, cyclin dependent kinase 2 (cdk 2), and cyclin dependent kinase 4 (cdk 4);~~

~~inactivation of the target protein can ameliorate a disease or pathology; and
the method comprises comprising the steps of:~~

(a) producing a library of comprising human protease muteins comprising mutation(s) of a human protease scaffold and/or biologically active portions thereof at an amino acid position selected from among amino acid residues between 58 and 64, 97, 98, 99, 100, 171, 174, 180, 189, 190, 191, 192, 215, 217, 218 and 226 by chymotrypsin numbering, wherein:

each different protease mutein in the library is a member of the library;

each member having N mutations relative to a wild-type human protease scaffold or a biologically active portion thereof, wherein N is a positive integer; and

the human protease scaffold is selected from among a granzyme A, granzyme B, granzyme M, cathepsin, MTSP-1, elastase, chymase, tryptase, chymotrypsin, collagenase, factor Xa, Protein C, plasma kallikrein, plasmin, trypsin, thrombin, complement factor serine proteases, papain, ADAMTS13, endopeptidase, furin, cruzain and urokinase plasminogen activator (uPA); and

(b) contacting members of the library with the target protein or with a polypeptide comprising a substrate sequence that is present in the target protein, wherein:

the target protein is selected from among a caspase 3, tumor necrosis factor, tumor necrosis factor receptor, interleukin-1, interleukin-1 receptor, interleukin-2, interleukin-2

receptor, interleukin-4, interleukin-4 receptor, interleukin-5, interleukin-5 receptor, interleukin-12, interleukin-12 receptor, interleukin-13, interleukin-13 receptor, p-selectin, p-selectin glycoprotein ligand, Substance P, Bradykinin, PSGL, factor IX, immunoglobulin E, immunoglobulin E receptor, CCR5, CXCR4, glycoprotein 120, glycoprotein 41, hemagglutinin, respiratory syncytium virus fusion protein, B7, CD28, CD2, CD3, CD4, CD40, vascular endothelial growth factor, VEGF receptor, fibroblast growth factor, endothelial growth factor, EGF receptor, TGF receptor, transforming growth factor, Her2, CCR1, CXCR3, CCR2, Src, Akt, Bcl-2, BCR-Abl, glucagon synthase kinase-3, cyclin dependent kinase-2 (cdk-2), and cyclin dependent kinase-4 (cdk-4); and
cleavage of a substrate sequence in the target protein inactivates an activity of the target protein;

(c) measuring a cleavage activity and/or substrate specificity of at least two members of the library for the substrate sequence or the target protein; and

(d) based on the measured activity and/or specificity, identifying those members of the library that have an increased cleavage activity and/or altered substrate specificity for cleaving the substrate sequence or the target protein relative to the wild-type human protease scaffold; and

(e) testing the identified protease(s) for cleavage and inactivation of an activity of the target protein that contains the substrate sequence, thereby identifying a protease mutein or a biologically active portion thereof that inactivates an activity of a target protein that is involved with the disease or pathology.

64. (Cancelled)

65. (Previously presented) The method of claim 63, wherein the human protease scaffold is selected from among granzyme A, granzyme B, granzyme M and MTSP-1.

66. (Currently amended) The method of claim 63 ~~any one of claims 63-65~~, wherein the target protein is selected from among caspase 3, vascular endothelial growth factor and VEGF receptor.

67. (Previously presented) The method of claim 1, wherein the substrate sequence is pre-selected so that its cleavage in the target protein inactivates the target protein.

68. (Previously presented) The method of claim 1, wherein the substrate sequence comprises a P4-P1 tetrapeptide substrate.

69. (Previously presented) The method of claim 1, wherein the target protein or substrate sequence is fluorogenically labeled.

70. (Previously presented) The method of claim 53, wherein the substrate sequence is pre-selected so that its cleavage in the target protein inactivates the target protein.

71. (Previously presented) The method of claim 53, wherein the substrate sequence comprises a P4-P1 tetrapeptide substrate.

72. (Previously presented) The method of claim 53, wherein the target protein or substrate sequence is fluorogenically labeled.

73. (Previously presented) The method of claim 59, wherein the substrate sequence is pre-selected so that its cleavage in the target protein inactivates the target protein.

74. (Previously presented) The method of claim 59, wherein the substrate sequence comprises a P4-P1 tetrapeptide substrate.

75. (Previously presented) The method of claim 59, wherein the target protein or substrate sequence is fluorogenically labeled.

76. (Previously presented) The method of claim 63, wherein the substrate sequence is pre-selected so that its cleavage in the target protein inactivates the target protein.

77. (Previously presented) The method of claim 63, wherein the substrate sequence comprises a P4-P1 tetrapeptide substrate.

78. (Previously presented) The method of claim 63, wherein the target protein or substrate sequence is fluorogenically labeled.